

ACCESSION NR: AP4019498

conductance decreases with a decrease of the dielectric constant of the solvent, except for nitrobenzenes in which the solubility and electric conductance are insignificant. Thus the electric conductance of  $\text{NbOCl}_3$  depends not only on the dielectric constant of the solvent but also on its chemical nature. The solubility in dioxane (dielectric constant = 2.18) is greater than in chloroform (dielectric constant = 4.98), and greater than in nitrobenzene and in acetone. The stability of methanol and dioxane solutions of  $\text{NbOCl}_3$  to hydrolysis is indicative of the formation of strong molecular compounds or solvates between  $\text{NbOCl}_3$  and the solvent. The dipole moment of  $\text{NbOCl}_3$  in dioxane is  $4.80 \pm 0.1D$ . The obtained data was interpreted from the standpoint of chemical interaction between  $\text{NbOCl}_3$  and the solvent, the oxygen of the alcohol or dioxane or the nitrogen of the pyridine acting as electron donor, and the niobium atom as the acceptor. A plot of the conductance of  $\text{NbOCl}_3$  in the alcohols as a function of temperature shows a straight line function with methanol. Curves for propanol, butanol and heptanol go through a maximum, the lower the alcohol radical and the higher the temperature at which the maximum conductance occurs. The dielectric constant, refractive index, density and molar polarization of various concentrations of  $\text{NbOCl}_3$  in dioxane are tabulated. Orig. art. has: 4 tables and 1 figure

Cord 2/3

ACCESSION: NR: AP4019498

ASSOCIATION: Rostovskiy-na-Donu gosudarstvennyy universitet (Rostov-on-Don  
State University)

SUBMITTED: 07Feb63 DATE ACQ: 31Mar64 ENCL: 00

SUB CODE: CH NO EXP Sov: 002 OTHER: 006

Cord 3/3

OSIPOV, O.A.; MINKIN, V.I.; KOGAN, V.A.

Dipole moments and the structure of  $\alpha$ -hydroxyaldehyde anils. Zavod. khim. 37 no. 1:1492-1499 Jl '63. MIRA 17:2)

1. Rostovskiy-na-Donu gosudarstvennyy universitet.

GAYVORONSKIY, V.I., OSIPOV, O.A., SHAGIDULLIN, R.R.

Infrared spectra of thorium, zirconium, and uranyl complex compounds  
with 8-oxin and tributyl phosphate. Radiokhimiia 5 no.2:244-248  
'63. (MIRA 16:1C)

DOROFEEVYENKO, G. N.; BABIN, Ye. P.; ROZENBERG, B. A.; OSIPOV, O. A.;  
KASHIRENINOV, O. Ye.

Catalytic acetylation of some polymers. Izv. vys. ucheb. zav.:  
khim. i khim. tekhn. 5 no.5:804-807 '62.  
(MIRA 16:1)

1. Donetskoye otdeleniye Instituta organicheskoy khimii AN  
UkrSSR i Rostovskiy-na-Donu gosudarstvennyy universitet.

(Polymers) (Acetylation)

OSIPOV, O.A.; SEMENOVA, I.M.

Infrared spectra of the molecular compounds of indium  
chlorides with nitrogen-containing organic substances.  
Zhur. ob. khim. 33 no. 3:720-724 Mr '63. (MIRA 16:3)

1. Rostovskiy-na-Donu gosudarstvennyy universitet.  
(Indium chloride)  
(Nitrogen compounds--Absorption spectra)

S/186/63/005/002/003/005  
E075/E136

AUTHORS: Gayvoronskiy, V.I., Osipov, O.A., and Shagidullin, R.R.

TITLE: Infrared spectra of complex compounds of thorium,  
zirconium, and uranyl with 8-hydroxyquinoline and  
tributylphosphate

PERIODICAL: Radiokhimiya, v.5, no.2, 1963, 244-248

TEXT: The authors investigated the structure of the complex compounds by studying infrared spectra of the solid internal complex compounds dispersed in vaseline. For Th and UO<sub>2</sub>, in addition to complexes Th (C<sub>9</sub>H<sub>6</sub>ON)<sub>4</sub> and UO<sub>2</sub>(C<sub>9</sub>H<sub>6</sub>ON)<sub>2</sub>, the complexes with the composition of 1:5 and 1:3 were also synthetized. The spectra of the complexes and those of 8-hydroxyquinoline in the frequency range 700 to 3000 cm<sup>-1</sup> indicate that the double bonds in the aromatic rings conjugate to an increased degree in the complexes. The conjugation is probably transmitted through the complexing metal. There are three strong absorption bands in the spectra of the complexes in the region of 400 - 700 cm<sup>-1</sup>, which are not present in the spectrum of 8-hydroxyquinoline. These bands

Card 1/2

Infrared spectra of complex ...

S/186/63/005/002/003/005  
E075/E136

shift towards longer wavelengths (from 485, 505 and 605 cm<sup>-1</sup> to 495, 515 and 615 cm<sup>-1</sup>) on passing from the heavy metals to Zr. Differences were also found between the absorption bands at 700-825 cm<sup>-1</sup> of the complexes due apparently to differences in their crystal structure. The strong absorption band for the UO<sub>2</sub> complex at 917 cm<sup>-1</sup>, absent in other complexes, is ascribed to the asymmetric vibration of UO<sub>2</sub> group. The spectra of the mixtures of the metal chlorides and tributylphosphate (TBP) indicate that a molecular compound between the metals and TBP is formed via the oxygen in P = O group.  
There are 3 figures and 2 tables.

SUBMITTED: July 5, 1961

Card 2/2

OSIPOV, O.A.; MINKIN, V.I.; KOGAN, V.A.

Effect of the chemical structure of organic complex-forming  
anions on the polarity of complex compounds. Zhur. fiz. khim.  
36 no.4:889-894 Ap '62. (MIRA 15:6)

1. Rostovskiy universitet.  
(Complex compounds)

S/137/62/000/012/065/085  
A006/A101

AUTHORS: Lesnykh, D. S., Lifshits, Ya. G., Osipov, O. A., Smovt, M. S.

TITLE: An electrochemical method of metal sulfonation

PERIODICAL: Referativnyy zhurnal, Metallurgiya, no. 12, 1962, 131 - 132,  
abstract 21808 ("Uch. zap. Rostovsk.-n/D. un-ta", 1959, v. 60,  
151 - 172)

TEXT: The authors studied the effect of factors upon the quality of a forming surface layer in the electrochemical method of cast-iron and steel sulfonation. These factors are: bath composition; metal type used for the cathode specimen (part) prior to sulfonation; current density on the anode; bath temperature; duration of the process and throwing power of the bath. Aqueous solutions and melts of sulfur-containing salts were used as sulfonation baths, e.g.  $\text{CH}_3\text{COOK}$  50%,  $\text{CH}_3\text{COONa}$  30%,  $\text{Na}_2\text{S}_2\text{O}_3$  10% and KCNS 10% with an operational temperature of the melt as high as 260 - 240°C, and a 10% aqueous solution of  $\text{Na}_2\text{S}_2\text{O}_3$  and KCNS in a 5 to 2 ratio; the sulfonation temperature is 50 - 75°C. To obtain a strong and elastic sulfonated layer, the specimens were

Card 1/2

OSIPOV, O.A.; MINKIN, V.I.; KASHIRENINOV, O.Ye.

Physicochemical properties of resins obtained by polycondensation  
of benzyl chloride and 1-chloromethylnaphthalene in the presence  
of chlorides of elements of the fourth group. Vysokom. soed. 3  
no.12:1774-1781 D (%) (MIRA 10:3)

1. Rostovskiy Gosudarstvennyy universitet.  
(Resins, Synthetic) (Toluene) (Naphthalene)

OSIPOV, O.A.; ARTEMIOVA, V.M.; KOGAN, V.A.; LYSENKO, Yu.A.

Dipole moments of the complex compounds of tin, titanium, and  
siliconium tetrachlorides with dibasic acid esters. Zhur. ob. khim.  
32 no.5:1368-1373 My '62. (MIRA 15:5)  
(Complex compounds—Dipole moments)

OSIPOV, O.A.; K. SHIRENINOV, Ye.

Interaction of  $TiCl_4$  and  $AlCl_3$  with organotin compounds. Zhur.-  
ob.khim. 32 no.6:17-1723 Je '62. (MIR: 15:6)

1. Rostovskiy-na-Donu gosudarstvennyy universitet.  
(Titanium chlorides) (Aluminum chloride)  
(Tin organic compounds)

MINKIN, V.I.; OSIPOV, O.A.; KOGAN, V.A.

Dipole moments and absorption spectra of o-hydroxyaldehyde anils.  
Dokl.AN SSSR 145 no.2:336-339 J1 '62. (MIRA 15:7)

1. Rostovskiy-na-Donu gosudarstvennyy universitet. Predstavлено  
akademikom B.A.Arbusovym.  
(Schiff Bases--Dipole moments)

OSIPOV, O.A.; PANINA, M.A.; KASHIRENINOV, O.Ye.; NEMIROV, G.V.;  
SHELOMOV, I.K.

Dielectric constant of binary liquid systems consisting of polar  
components. Zhur.ob.khim. 31 no.10:3153-3160 0 '61.  
(MIRA 14:10)

(Systems (Chemistry)) (Dielectrics)

OSIPOV, O.A.

Relation between the dielectric constant and mean dipole moments  
in mixtures of polar liquids. Uch.zap. RGU 41:3-25 '58.  
(KIRK 15:1)  
(Dielectric constants) (Liquids--Dipole moments)

OSIPOV, O.A.

Dipole moments in binaries of elements of the 4th group of the  
D.I.Mendeleev periodic system. Ucn.zap. MGU no.60:13-36 '59.  
(MLA L4:10)

(Complex) (Compounds- Dipole moments)

"APPROVED FOR RELEASE: Wednesday, June 21, 2000

CIA-RDP86-00513R001238

LESNYKH, D.S.; LIFSHITS, Ya.G.; OSIPOV, O.A.; SMOVT, M.S.

Electrochemical method for the sulfidization of metals. Uch.zap.  
KGU no.60:151-172 '59. (KTA 14:10)  
(Metals) (Sulfurization)

APPROVED FOR RELEASE: Wednesday, June 21, 2000

CIA-RDP86-00513R001238

KOVALENKO, P.N.; BAGDASAROV, K.N.; OSIFOV, O.A., dots., otv. red.;  
SHKORINOV, V.P., red.; PAVLICHENKO, M.I., tekhn. red.

[Physicochemical methods of analysis; practical handbook] Fiziko-  
khimicheskie metody analiza; prakticheskoe rukovodstvo. Rostov-na-  
Donu, Izd-vo Rostovskogo univ., 1962. 349 p. (MIRA 15:6)  
(Chemistry, Analytical) (Electrochemical analysis)

S/079/62/032/006/001/006  
D202/D304

AUTHORS: Osipov, O. A. and Kashireninov, O. Ye.

TITLE: Interactions of  $TiCl_4$  and  $AlCl_3$  with organic com-  
pounds of tin

PERIODICAL: Zhurnal obshchey khimii, v. 32, no. 6, 1962, 1717-  
1723

TEXT: Reactions of  $Sn(C_2H_5)_4$ ,  $Sn(C_2H_5)_3Cl$ ,  $Sn(C_2H_5)_2Cl_2$  and  
 $Sn(C_2H_5)Cl_3$  with  $TiCl_4$ ,  $ZrCl_4$ ,  $ThCl_4$  and  $AlCl_3$  were carried out to  
investigate the structures and catalytic properties of the re-  
sulting complexes. All these compounds react violently when mixed  
directly with  $TiCl_4$ . Only the first two give violent reactions  
with  $AlCl_3$ ; none of them reacts with  $ThCl_4$  and  $ZrCl_4$ . The reac-  
tions with  $TiCl_4$  and  $AlCl_4$ , when carried out in benzene, proceed  
smoothly without the formation of resins to give stable bimetallic

Card 1/2

OSILOV, O.A.; KLETENIK, Yu.B.

Reaction of zirconium chloride with organic additives having  
various functional groups. Zhur. ob. khim. 31 no.8:2451-2456 Ag  
'61. (MIRA 14:8)

(Zirconium chloride)  
(Zirconium organic compounds)

1369  
S 081 61 000 C. 175  
B17-B1

1800 2408

AUTHORS Kovalenko, E. N., Rozin, G. N., Osipov, V. A.  
Yevstafev, M. M., Kravtsov, Ye. Ye.

TITLE Anodizing in the presence of chloride ions, and the  
quality control of oxide films on the alloy D16T

PERIODICAL Referativnyy zhurnal. Khimika, no. 12, 1971, p. 2271-2274  
2<sup>2</sup>K144 "Fiz.-khim. metodi analiza i kontrolya"  
Rostov-na-Donu, Rost-VSK "Nauka", 1971.

TEXT An investigation is made of the effect of the presence of  
0.1 g/liter, in the tank, on the potential, depth of oxide film and the  
test time in the alloy D16T in the process of anodizing in a tank. It is  
found that it is possible to improve the potential of the anodizing bath by adding  
more than 0.1 g/liter chloride ions without affecting the depth or rate of anodizing.  
It is suggested that sheet D16T Duralumin should be anodized at a current density  
of 0.2 A/dm<sup>2</sup> in a solution containing 0.1 g/liter Cl<sup>-</sup>. Optimum conditions of anodizing  
with chloride ions have been found to be 0.2 A/dm<sup>2</sup> and 10 min.  
[Abstracted from "Chistoetchnaya metalloobrabotka", No. 1, 1971]

Cards 1

3/13/87  
A-1

AUTHORS: K. Valenkov, P. N. Rulin, G. M. Slipov, A. V. Yermakova, M. N. Kravtsov, Ye. Ye.

TITLE: Filling and control of anodized alloy Al-T (V.T) in the presence of bisulfite and sulfate ions.

PERIODICAL: Referativnyy zhurnal. Metalurgiya, no. 11, 1981, p. 1674-1676.  
USSR. (Vestn. "Fiz.-Khim. metody analiza i kontrolya proizvdenii"  
Rostov-na-Donu, 1981, no. 1, p. 13-16.)

TEXT: The authors studied the effect of the presence of bisulfite and sulfate ions upon the process of chromate filling of the oxide film on the D17 alloy. The dependence of the film quality (impurity and thickness of the film) upon the concentration of impurity ions is established. Sulfate ions suppress the chromate ion adsorption, as result of which the films have a lighter tint. It is recommended that films formed at high D be subjected to a longer filling. It is entirely possible to raise the admissible limit of admixtures in the filling vat from 1.5 to 3, and from 3 to 5 grams per liter for bisulfite and sulfates respectively. There are 6 references.

[Abstracter's note: Complete translation] Ye. Laymer  
Card 17

LYSENKO, Yu.A.; OSIFOV, O.A.

Interaction between titanium tetrachloride and chloromethyl acetate and ethyl stearate. Zhur. neorg. khim. 6 no.7:  
1656-1661 Jl '61.  
(MIRA 14:7)

1. Kubanskiy sel'skokhozyaystvennyy institut, kafedra  
organicheskoy fizicheskoy i kolloidnoy khimii.  
(Titanium chloride) (Esters)

15.8063

15.8063

S/190/61 604.072-11-1  
B1C1 B110

AUTHORS: Osipov, O. A., Mirkin, V. I., Kashirchenko, G. Ye.

TITLE: Physicochemical properties of resins obtained by polycondensation of benzyl chloride or 1-chloromethyl naphthalene with chlorides of elements of the fourth group

PERIODICAL: Vysokomolekulyarnyye soyedineniya, v. 1, n. 1, 1974  
1974 1981

TEXT: The purpose of this study was (1) to obtain data on the activity of halides of elements of the fourth group ( $\text{SiCl}_4$ ,  $\text{TiCl}_4$ ,  $\text{GeCl}_4$ ,  $\text{ZrCl}_4$ , and  $\text{ThCl}_4$ ); (2) to compare the polyphenylene methyl resins obtained by polycondensation of benzyl chloride with the polynaphthalene methyl resins obtained by polycondensation of 1-chloromethyl naphthalene. Polycondensation of benzyl chloride was achieved at a molar ratio catalyst: benzyl chloride = 1 : 50. The yield was 80 - 90%. Dark-red brittle substances were obtained, easily soluble in benzene, toluene, tetralin, carbon tetrachloride, and hexane, poorly soluble in ethanol, acetone, and other polar

Card 1/86

S, 190/61.005.012 000  
B'01, B'110

Physicochemical properties

solvents. The solutions fluoresced slightly. For the catalysts used it is indicated: with  $TiCl_4$  the reaction started at 20°C softening point; the resin 80°C, molecular weight (MW) 2100;  $SnCl_4$ , reaction started at 20°C, softening point 74°C, MW 2200;  $ZrCl_4$ , reaction started at 40°C softening point 74°C, MW 1800. With  $SiCl_4$  and  $TnCl_4$  no reaction was observed even after long heating at 100°C. Polycondensation of 1-chloromethyl naphthalene was carried out at equal ratio catalyst : monomer. Results are given in Table 2. The catalytic activity increased in the order:  $SiCl_4$  -  $TnCl_4$  -  $ZrCl_4$  -  $G_2Cl_4$  -  $SnCl_4$  -  $TiCl_4$ . The decrease of catalytic activity in the order  $TiCl_4$  -  $G_2Cl_4$  is due to the increasing of interaction between metal and chlorine. Highly soluble polymers were obtained at elevated temperatures. Thus, a significant increase of viscosity in tetrahydrofuran. The dielectric constant of the polymer was measured in tetrahydrofuran by a method described earlier Osipov et al. Zh. obshch. khim. 32, 16 (1956) (Table 1).  $\epsilon_{200^{\circ}C}$  is ascribed to infrared absorption. In solid phase  $\epsilon_{200^{\circ}C}$  was between 2.1 (polystyrene) methyl and 2.7

Card 2/1

Phyicochemical properties...

S/190/61/CU3/012/002/012  
B101/B110

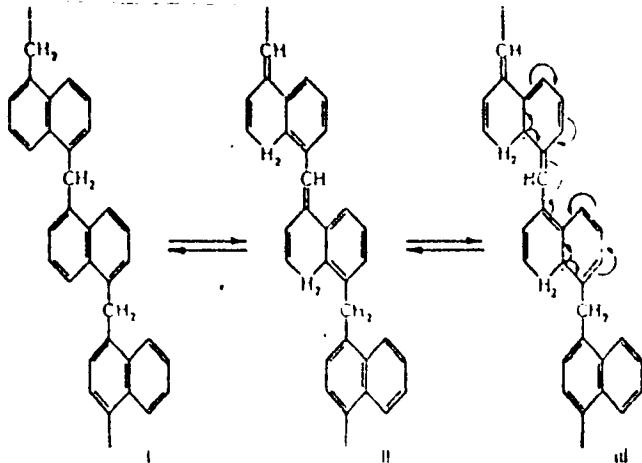
(polynaphthalene methyl). The polymers are recommended for impregnation of paper condensers. The magnetic susceptibility was determined according to L. G. Gouy.  $\pi \cdot 10^{-6} \text{ cm}^3/\text{g}$  for polynaphthalene methyl obtained with  $\text{SiCl}_4$  was: +1.441; with  $\text{TiCl}_4$ : -0.045; with  $\text{GeCl}_4$ : +0.892; with  $\text{ZrCl}_4$ : +0.226; with  $\text{SnCl}_4$ : +0.758; with  $\text{ThCl}_4$ : +0.331; with  $\text{TiCl}_4$  in hexane: +0.555; with  $\text{TiCl}_4$  in  $\text{CCl}_4$ : +0.130. Since polynaphthalene methyl contains no polar groups, its paramagnetism is ascribed to formation of free radicals in the macromolecules.

X

Card 3/ 6

50906  
S/190/61/003/012/002/012  
B101/B110

Physicochemical properties...



Both the polyphenylene methyl and the polynaphthylene methyl polymers form films from benzene solution which firmly adhere to the metal and protect it from corrosion. A small iron plate covered with polynaphthylene methyl

Card 4/15

OSIPOV, O.A.; KASHIRENINOV, O.Ye.

Magnetic susceptibility of some tin molecular compounds. Part 1.  
Zhur. ob. khim. 31 no.6:1755-1759 Je '61. (MIRA 14:6)

1. Rostovskiy-na-Donu gosudarstvennyy universitet.  
(Tin organic compounds—Magnetic properties)

S-079-01-001-014  
D228 D305

AUTHOR:

Kashchenko, O. V., USPENSKIY, O. A., Putilov, M. S.  
and Smirchenko, V. N.

TITLE:

Magnetic susceptibility of binary liquid systems

PUBLICATION:

Zhurnal Russicheskikh Khimicheskikh Nauk, 1964, 35(4), 1000

TEXT: The authors determined the magnetic susceptibility of binary liquid systems consisting of benzene and either ethyl benzoate (I), propylene glycol (II), caproic acid (III) or caproate (IV). The effect of polar solvents (I-IV) and nonpolar solvents (V-VII) on the magnetic susceptibility of benzene was also studied. The authors also studied the influence of the polarity of components (V-VII) on the magnitude of the magnetic susceptibility of mixtures. The work of the field suggests that there is a direct connection between the magnetic susceptibility of binary liquid systems and the influence of the components and of the dissimilarity of the magnetic susceptibility of components.

Card 1 of 3

Magnetic susceptibility

S-079-61-031-C11-197  
D228-B305

the additivity is greatest in systems consisting of polar components. Experimental procedure. All materials were first purified by the Auer-Schmidt method (cf. [16]). The glassy aluminum (96.32% Al) was obtained from the VEB Metallwerke Schwerin (DDR) in 1957. The samples of metallic salts were kindly furnished by the Institute of Inorganic Chemistry of the University of Berlin. The magnets were fitted to the working devices to maintain the field strength constancy and to eliminate convection currents. The apparent changes in the magnetic susceptibilities were measured by means of microbalance weights, and the calibrating material was purified water saturated with air with a magnetic susceptibility of  $\chi = 0.293 \times 10^{-6}$ . Experimental suspensions and their dilutions. The data show that the magnetic susceptibility of the systems I - IV whose components possess intrinsic dipole moments follows a rectilinear course at the low concentration levels studied by the authors. For other systems, where the components have chemically strong interactions, the components' dipole moments are more important, since the behavior is largely determined by the behavior of the components themselves.

Carlo S.

Magnetic susceptibility

S-679 61-031-C14-00000  
B228 D305

With the exception of system V, the deviations of the isotherms from the rectilinear course have positive values, and the maximum deviations correspond to the composition of the resulting compound. The authors consider that magnetic susceptibility may find a wide application in physico-chemical analysis. There are 10 tables and 26 references, 12 Soviet bloc and 16 non-Soviet bloc. The references to the 4 most recent English-language publications read as follows: 1. van Vleck, *The Theory of Electric and Magnetic Susceptibilities*, Oxford University Press, 1932; 2. Trew, D. & Watkins, *Trans. Faraday Soc.*, 29, 310 (1933); 3. Seely, *Phys. Rev.*, 51, 812 (1936); 4. Morris, D. *Philosophical Magazine*, 36, 101 (1947).

SUBMITTER:

Card 3/3

"APPROVED FOR RELEASE: Wednesday, June 21, 2000

CIA-RDP86-00513R001238

KASHIRENIKOV, O.Ye.; OSIPOV, O.A.; PANINA, M.A.; MARCHENKO, V.N.

Magnetic susceptibility of binary liquid systems. Zhur. ob. khim.  
31 no. 11:3504-3509 N '61.  
(ZIFK 14, 11)  
(Systems (Chemistry)--Magnetic properties)

APPROVED FOR RELEASE: Wednesday, June 21, 2000

CIA-RDP86-00513R001238

OSIPOV, O.A.; SIMONOV, A.M.; MINKIN, V.I.; GARNOVSKIY, A.D.

Dipole moments of imidazole and its derivatives. Dokl.AN SSSR 137  
no.6:1374-1376 Ap '61. (MIRA 14:4)

1. Rostovskiy-na-Donu gosudarstvennyy universitet. Predstavлено  
академиком М.М.Shemyakiny. (Imidazole—Dipole moment\*)

KLETENIK, Yu.B.; OSIPOV, O.A.

Reaction of titanium tetrahalides with esters of acetic acid.  
Zhur. ob. khim. 31 no.3:710-716 Mr '61. (MIRA 14:3)

1. Khimiko-metallurgicheskiy institut Sibirskogo otdeleniyu AN SSSR.  
(Titanium compounds)

OSIPOV, O.A. ; LYSENKO, Yu.A.

Electric properties of systems formed by titanium tetrachloride  
with esters of trichloroacetic acid. Zmr. ob. khim. 30 no.12; 3866-  
3869 D '60. (MIRA 13:12)

1. Rostov-na-Donu gosudarstvennyy universitet.  
(Titanium chloride) (Acetic acid)

OSIPOV, O.A.; KLETENIK, Yu.B.

Heats of reactions between zirconium halides and esters. Zhur. neorg.  
khim. 5 no.10:2220-2222 O '60. (MIRA 13:10)  
(Zirconium halides) (Chemical reaction, Heat of)

3/673/60/050, 012, 004 D  
B001/B064

AUTHORS: Osipov, O. A. and Lysenko, Yu. A.  
TITLE: Electrical Properties of the Systems Formed From Titanium  
Tetrachloride and Esters of Trichloro Acetic Acid  
PERIODICAL: Zhurnal obshchey khimii, 1960, Vol. 30, No. 1,  
pp. 3866-3869

TEXT: This paper gives the dielectric constants of the systems of  $TiCl_4$  with ethyl trichloro acetate, n-butyl trichloro acetate, isobutyl trichloro acetate, isotrichloro amyl acetate, and the dipole moments of some etherates  $TiCl_4 \cdot E$ , where  $E$  is the molecule of the ester. All determinations were carried out by previous methods (Refs. 1, 3, 5) at  $20 \pm 0.1^\circ C$ . Tables 1-4 show the dielectric constants ( $\epsilon$ ), densities ( $d$ ), refractive indices  $n_D^2$ , as well as the values of the orientation polarization and the mean dipole moments of the systems of  $TiCl_4$  and the four esters of trichloro acetic acid. The two last columns of the table show the deviations of the dielectric constants and polarization from the additive values.

Card 1/3

Electrical Properties of the Aluminothermic S, 279, 60, 275, 512, 501  
From Titanium Tetrachloride and Esters of Tri- B001/B064  
chloro Acetic Acid

members of a homologue series (Tables 1-4). There are 11 references and  
11 references: 9 Soviet, 1 US, and 1 British.

ASSOCIATION: Rostovskiy-na-Donu gosudarstvennyy universitet  
(Rostov-na-Donu State University)

SUBMITTED: February 1, 1961

Card 3/3

53700 2209 1236 1273

86494  
S'078/60 1236 1273 XX  
BC23/BC44

AUTHORS: Osipov, O. A., Lysenko, Yu. A.

TITLE: Heat of Formation of Esterates of Titanium Tetrachloride

PERIODICAL: Zhurnal neorganicheskoy khimii, 1960, Vol. 5, No. 8,  
pp. 1840-1845

TEXT: The authors investigated the heats of mixing of the systems:

 $TiCl_4$  - iso- $CH_3COOC_5H_{11}$ ;  $TiCl_4$ -sec- $CH_3COOC_8H_{11}$ ; $TiCl_4$  - n- $HCOOC_4H_9$ ; and  $TiCl_4$  - iso- $HCOOC_5H_{11}$ .

On the basis of the results obtained per g-mole, also the chemical nature of each component may be determined. For systems in which chemical processes take place, the interaction between the components is clearly seen if the thermal effects per g-mole of each component are calculated (Figs. 1-4, Tables 1-4). Further investigations of the authors confirm the hypothesis according to which compounds of the type  $2TiCl_4 \cdot E$  ( $E$  = ester) are not formed in the liquid phase. It was shown that in the compounds  $TiCl_4 \cdot E$ ,

Card 1/6

Heat of Formation of Esterates of Titanium  
Tetrachloride

86494  
S'078/60/065/5 8 1 15/11/XX  
B023/B066

titanium has the coordination number 5, with the ester oxygen of the ester molecule being the principal electron donor. There are 4 figures, 5 tables, and 11 references: 9 Soviet and 2 German.

SUBMITTED: March 31, 1959

Legend to Fig. 1: Heats of mixing in the system  $TiCl_4 - i-C_5H_9OCCH_3$ , kcal/mole; 1: thermal effects of  $TiCl_4$ ; 2: thermal effects of the ester;

3: thermal effects of the mixture.

Legend to Fig. 2: Heats of mixing in the system  $TiCl_4 - sec-C_6H_{11}OCCH_3$ ,

kcal/mole; for 1,2,3 see Fig. 1.

Legend to Fig. 3: Heats of mixing in the system  $TiCl_4 - n-C_4H_9OCCH_3$ ,

kcal/mole; for 1,2,3 see Fig. 1.

Legend to Fig. 4: Heats of mixing in the system  $TiCl_4 - i-C_6H_{11}^+CH_3$ ,

kcal/mole; for 1,2,3 see Fig. 1.

Card 2/6

86494  
 S/078/60/005/008/070/071/XX  
 B023/PC66

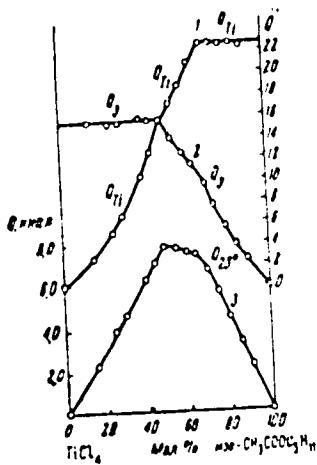


Рис. 1. Термоплоты смешения системы  $\text{TiCl}_4$  — изо- $\text{C}_6\text{H}_4\text{OOCCH}_3$ , ккал/г-моль

1 — тепловые эффекты четыреххлористого титана; 2 — тепловые эффекты эфира; 3 — тепловые эффекты смеси

Card 3/6

86494  
S/078/60/005/008/C30/031/XX  
B023/B066

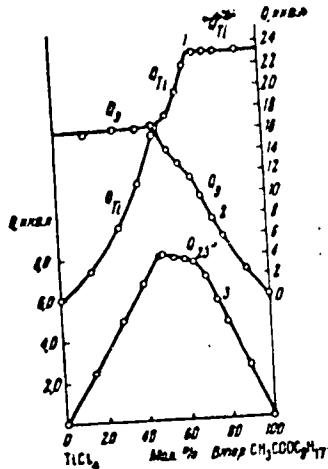


Рис. 2. Теплоты смешения системы  $TiCl_4$  — втор.- $C_4H_7OOCCH_3$ , ккал/г-моль.

1 — тепловые эффекты четыреххлористого титана; 2 — тепловые эффекты вина; 3 — тепловые эффекты смеси

Card 4/6

86b94  
 S/078/F0/005/008/070/031/XX  
 B023/B060

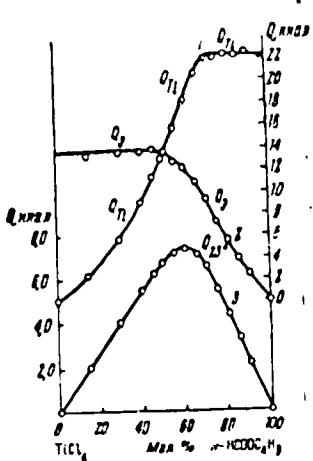


Рис. 3. Теплоты смешения системы  
 $TiCl_4-n-C_4H_9OSi$ , ккал/с-моль.  
 1 — тепловые эффекты четыреххлористого титана; 2 — тепловые эффекты  
 «фира»; 3 — тепловые эффекты смеси

Card 5/6

86194

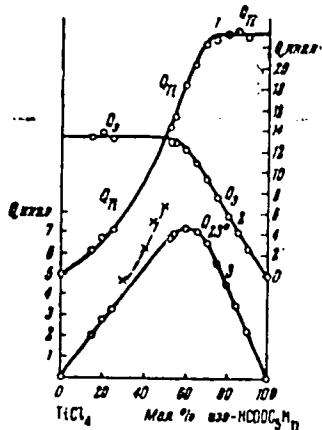
S/078/60/005/006/030/031/xx  
B023/B066

Рис. 4. Термодинамика смеси системы  
 $TiCl_4$ —изо- $C_4H_{11}OOCH_3$ .  
ккал/моль.

1 — тепловые эффекты четыреххлористого титана; 2 — тепловые эффекты эфира;  
3 — тепловые эффекты смеси

Card 6/6

S/078, 60/005/0'0/0<sup>4</sup>'0'0 XX  
B017, B06

AUTHORS: Osipov, O. A. and Kletenik, Yu. B.  
TITLE: The Heat of the Reaction of Zirconium Halides With Esters  
PERIODICAL: Zhurnal neorganicheskoy khimii, 1960, Vol. 5, No. 10,  
pp. 2220-2222

TEXT: The heat which is generated in the reaction of zirconium bromide and zirconium chloride with ethyl formate, ethyl acetate, and ethyl butyrate, as well as the heat of solution of zirconium iodide in ethyl acetate were determined. The ratio of zirconium bromide and zirconium chloride to the esters was 1 : 1, and 1 : 2, respectively. The heats of solution of zirconium halides and their complex compounds in these esters which were also determined are given in Table 1. Table 2 shows the heat of formation of the complexes. The complexes could be obtained only as solutions, not in the solid form. The heat of formation of complexes with the composition 1 : 1 was, in all cases investigated, higher than that of complexes with the composition 1 : 2. This result confirms the

Card 1/2

The Heat of the Reaction of Zirconium  
Halides With Esters

S/078/60/005, 010, 023, 030/XX  
B017/B067

dissociation scheme suggested earlier by the authors (Refn. 4,5). In contrast to molecular compounds of tetrahalides of tin in the order chloride - bromide - iodide with these esters, no continuous decrease in the heats of formation in the above order was observed in the complexes of zirconium halides. The heat of formation of complexes of zirconium bromide with these esters is somewhat higher than that of complexes of zirconium chloride. The heat of solution of zirconium iodide in ethyl acetate is 4% lower than the heat of solution of zirconium chloride in the same ester. There are 2 tables and 9 references: 7 Soviet and 2 US.

SUBMITTED: July 3, 1959

Card 2/2

KLETENIK, Yu.B., OSIPOV, O.A.

Physicochemical investigation of some zirconium halide complexes  
with esters of monobasic carboxylic acids. Izv.vys.ucheb.zav.;  
khim.i khim.tekh. 2 no.5:679-684 '59. (MIR 13:8)

1. Rostovskiy-na-Donu gosudarstvennyy universitet, kafedra  
fizicheskoy i kolloidnoy khimii.  
(Zirconium compounds)

OSIPOV, O.A.; PANINA, M.A.; YAGUBYAN, Ye.S.

Reacts of mixing dioxane with chloroform and *O*-toluidine.  
Zhur. ob. khim. 30 no.7:2127-2130 J1 '60.  
(MIRA 13:7)

1. Rostovskiy-na-Donu gosudarstvennyy universitet.  
(Dioxane) (Chloroform) (Toluidine)

OSIPOV, O.A.; ANTRONOVA, V.M.

Electric properties of inner complex compounds. Dokl. AN SSSR  
133 no.1:166-169 J1 '60. (MIRA 13:7)

1. Rostovskiy-na-Donu gosudarstvennyy universitet. Predstavleno  
akademikom A.N. Frumkinyem.  
(Complex compounds--Electric properties)

AUTHORS: Osipov O.A. Lyashko Yu A

78-3723/44

TITLE: XI The Electrolysis of Tetrachlorotitanium Compounds With Some Esters of Monohydric Acids (XI. Elektroliz soyedineniy chetyrekhkhloridogo titana s nekotorym estirami odnoosnovnykt. kislot)

PERIODICAL: Zhurnal neorganicheskoy khimii, 1958, v. 1, N 7, pp. 1605-1607 (USSR)

ABSTRACT: In the course of the present paper the results obtained by the electrolysis of the solutions of titanium(IV)-chloride in n-butyl-formate, isobutylacetate, ethyl acetate, and ethyl formate are investigated. Graphite and in some cases, platinum and silver were used as materials for electrodes. In the course of electrolysis there is a black precipitation at the cathode in the case of all experiments in which the ratio between titanium and chlorine is 1 : 3. At the anode a product is separated in the case of which the ratio between titanium and chlorine is 1 : 4. On the strength of these investigations it is assumed that at the cathode  $TiCl_3$  is first separated which is reduced with  $TiCl_4$  to  $TiCl_2$ . The formation of  $TiCl_2$  is characterized by the fact that at first

Card 1/2

**XI. The Electrolysis of Tetrachlorotitanium Compounds  
With Some Esters of Monobasic Acids** SJV 76-3 7-23/44

a yellowish brown precipitation is formed on the cathode. The results obtained confirm the scheme of the electrolytic dissociation of titanium-(IV) chloride in complex esters as described in earlier works. There are 1 table and 9 references, 8 of which are Soviet.

**SUBMITTED** June 11, 1967

1. Titanium compounds--Electrolysis
2. Titanium compounds--Separation
3. Titanium compounds--Precipitation
4. Titanium compounds--Test results

Card 2/2

81731  
S/020/60/133/01/46/070  
B004/B007

53700(A)  
AUTHORS: Osipov, O. A., Artemova, V. M.

TITLE: The Electrical Properties of Intracomplex Compounds<sup>1</sup>

PERIODICAL: Doklady Akademii nauk SSSR, 1960, Vol. 133, No. 1,  
pp. 166 - 169

TEXT: The authors investigated the electrical properties of the acetyl acetones and hydroxyquinolates of zirconium, thorium, indium, beryllium, chromium, copper and cadmium, which have a chelate bond. The present investigation aimed at ascertaining the cause of the great difference between molecular and electron polarization. The dielectric constant, the density, and the refractive index in solutions (carbon tetrachloride, benzene, dioxane) were determined. Table 1 gives the values found, which show that the dipole moments do not depend on the nature of the solvent, and that the difference between molecular and electron polarization is not caused by this solvent. The authors assumed the cause to be a shifting of the atomic nuclei due to the application of a voltage, and therefore investigated the electrical properties of these compounds in the solid phase. ✓

Card 1/2

OSIPOV, O.A.

Deposition potentials of zinc and hydrogen in the presence of potassium  
xanthogenate. Uch. zap. RGU 40:103-1.1 '58. (MIRA 13:10)  
(Xanthic acid) Zinc  
(Hydrogen) (Reduction, Electrolytic)

OSIPOV, C.A.; MINKIN, V.I.; TUMAKOVA, Zh.A.

Dipole moments and structure of bis-salicylalacetylimidazoles (part II). Zhur. strukt. khim. 5 no.6(918-919) N.D. 1974. MIRAN 1975.

I. R. stroyekovskogo na gosudarstvennyy universitet.

"APPROVED FOR RELEASE: Wednesday, June 21, 2000

CIA-RDP86-00513R001238

MINKIN, V.I., GORELOV, M.I., CHIBOV, O.A.; OSTROUMOV, Yu.A.

Electron structure and absorption spectra of salicyialanilides  
and its derivatives. Opt. i spektrosc. 18 no.4:571-578 Ap '65.  
(MIRA 18:8)

APPROVED FOR RELEASE: Wednesday, June 21, 2000

CIA-RDP86-00513R001238

ISMAYLOV, Kh.M.; OSIPOV, V.A.; GADJIEV, M. I. et al. Neftegaz. i gornye promstv., 1976, No. 10, p. 10.  
CHIKINA, N.I.

Complex compounds containing phenol with sulfur and thiomethyl groups  
phenols and their derivatives. In: K. AN Azer. Nauk. Rada, no. 313, 1972  
1975.

1. Institut neftegaz. i gornye promstv. na. Ya. N. Mamedova, Baku,  
AN Azerb. SSR. 2. Rostovskiy gos. univ. im. N. G. Chernysheva, Rostov-na-Donu.

"APPROVED FOR RELEASE: Wednesday, June 21, 2000

CIA-RDP86-00513R001238

KOPI T. WY, TURK, 1961, 1962, 1963, 1964

Reproduction of this document is authorized only if it is determined that it is required for  
the conduct of intelligence operations.

It is the intent of the CIA to declassify this document by January 1, 2000.

APPROVED FOR RELEASE: Wednesday, June 21, 2000

CIA-RDP86-00513R001238

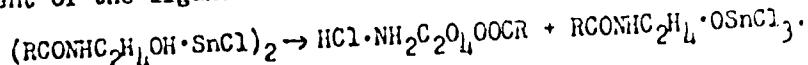
L 11/09-67 S.T(m)/SNT(j) RM  
Acc No: A77003666

SOURCE CODE: UR/0079/66/036/003/1484/1487

KOZLOV, I.A., OSIPOV, O.A., Volgodonskaya Affiliate, All-Union Scientific Research Institute of Synthetic Oil Substitutes (Volgodonskoy filial Vsesoyuznogo nauchno-issledovatel'skogo instituta sinteticheskikh zhirozameniteley) "Reaction of Tin and Titanium Tetrachlorides with Ethanolamides of Aliphatic Acids. III"

Moscow, Zhurnal Obshchey Khimii, Vol 35, No 8, 1966, pp 1484-1487

Abstract: It was found that in the reaction of tin and titanium tetrachlorides with monoethanolamides (MEA)  $\text{RCOHC}_2\text{H}_4\text{OH}$  and diethanolamides (DEA)  $\text{RCO}(\text{C}_2\text{H}_4\text{OH})_2$  of aliphatic acids, where R is a hydrocarbon radical containing from seven to 11 carbon atoms, complex compounds with the composition  $2\text{MEA} \cdot 2\text{MeCl}_4$  and  $2\text{DEA} \cdot 3\text{MeCl}_4$  are formed. It was further found that in dioxane solution these complex compounds undergo further molecular transformations: a molecular rearrangement of complex compounds of tin and titanium tetrachlorides with ethanolamides of aliphatic acids was detected and studied. A mechanism was proposed for the rearrangement: when solutions of the complexes in dioxane are heated, there is a replacement of the hydrogen of the hydroxyl group of the ethanolamides by the  $\text{SnCl}_3$  or  $\text{TiCl}_3$  group, liberating a  $\text{HCl}$  molecule. The latter causes a rearrangement of the ligand molecule, according to the scheme:



UDC: 547.258.11 + 547.372

Card 1/2

ACC NR: AP7005109

SOURCE CODE: UR/0079/66/036/009/1693/1702

KOLODYAZHNYI, Yu. V., MARCHENKO, V. N., OSIPOV, O. A., KOGAN, M. G., Rostov-on-Don State University (rostovskiy-na-Donu gosudarstvennyi universitet)

"Investigation of the Interaction Between Tetra-n.-Butoxytitanium and the Tetrachlorides of Tin and Silicon"

Moscow, Zhurnal Obshchey Khimii, Vol 36, No 9, 66, pp 1693-1702

Abstract: With the aid of various physicochemical techniques (dielectric losses, cryoscopy, electric conductivity, etc.) it is shown that tetrabutoxytitanium  $Ti(OBu)_4$  forms conducting complex compounds not only with stannic tetrachloride but also with such a weak electron acceptor such as silicon tetrachloride. It was shown that the interaction between the tetrachlorides of tin and silicon and tetra-n-butoxytitanium in dilute benzene solutions leads to the formation of the following complexes:  $SiCl_4 \cdot Ti(OBu)_4$ ,  $SiCl_4 \cdot 2Ti(OBu)_4$ ,  $SiCl_4 \cdot [Ti(OBu)_4]$ ,  $SiCl_4 \cdot [Ti(OBu)_4]^+$ . The association of complexes 1:2 composition was established and this is attributed to not only donor-acceptor interaction between molecules of tetrabutoxytitanium but also, and to a large degree, to the interaction between the butoxy-group hydrocarbon radicals; the gradual decomposition of such associated complexes accounts for the decrease in their electric conductivity with time. Orig. art. has 11 figures, 2 formulas and 8 tables. (JPIS: 38, 970)

TOPIC TAGS: organotitanium compound, organotin compound, organosilicon compound

SUB CODE:07 / SUBM DATE: 06Jul65 / ORIG REF: 013 / OTH REF: 001

Card 1/1

UDC: 547.1'3 + 546.81 + 546.28

OSIPOV, C. P. I. SHITENYAR, L. V.

Fertilizers and cultures

Effect of mineral fertilizers on hay crop in Niperne-Karabakh, C. P. Osipov, L. V. Shitennyar, Tern. baza, 2, No. 12, 1951

Q. Monthly List of Russian Accessions, Library of Congress, Hay 1954, Uncl.

CHIRCHIN, C. I., CHIRCHIN.

Crusader

Effect of mineral fertilizers on soy crop in Baranov-Karakakh., Farm. taza, 7, no. 12, 1951

9. Monthly List of Russian Accessions, Library of Congress, May 1951, 2 vols.

"APPROVED FOR RELEASE: Wednesday, June 21, 2000

CIA-RDP86-00513R001238

9. Monthly List of Russian Accessions, Library of Congress, Washington, D.C., 1962.

APPROVED FOR RELEASE: Wednesday, June 21, 2000

CIA-RDP86-00513R001238

MAMEDOV, Sh.A.; OSIPOV, O.B.; MAMEDOVA, A.R.

"Etheran"- a new type of agricultural poison [in Azerbaijani with  
summary in Russian]. Izv. AN Azerb. SSR. Ser. fiz.-tekhn. i khim.  
nauk no.1:125-131 '59. (MIRA 12:6)  
(Ether) (Insecticides)

MAMEDOV, Shamkhal; OSIPOV, O.B.; ALIYEVA, Kh.M.

Catalytic alkoxyethylchlorination of olefins. Azerb.khim.  
zhur. no.2:83-91 '60. (MIRA 14:8)  
(Olefins) (Ether)

NAMEDOV, Shamkhal; MAGERRAMOV, B.G.; OSIROV, O.B.; ALESKEROV, A.S.

Bactericidal properties of certain ether preparations. Azerb.  
khim.zhur. no.1:65-69 '61. (MIRA 14:8.  
(Ether) (Bactericides)

MAMFICOV, Shamhal; OSIPOV, C.B.; ALIYEVA, Kh.M.; ZEYNALOVA, V.M.

Efiran-66, a new herbicide. Dokl.AN Azerb.SSR 17 no.4:331-334  
'61. (MIRA 14:6)

1. Institut neftekhimicheskikh protsessov AN AzerSSR.  
Predstavлено академиком AN Azerbaydzhanskoy SSR V.R. Volobuyevym.  
(Herbicides) (Isopropyl ether)

MAMEDOV, Shamxhal; OSIPOV, O.B.; ALIYEVA, Kh.M.

"Betaefiran" (ether preparation). Azerb.khim.zhur. no.5:59-63  
'61. (MIRA 15:5)  
(Ethers) (Insecticides)

Shamkhal; O.B.; GRISHINA, Ye.N.

Efiran-59 and Efiran-94, new highly effective chemicals for  
bollworm control. Dokl. AN Azerb. SSR 17 no.8:691-695 '61.  
(MIRA 14:10)

1. Institut neftekhimicheskikh protessov AN AzerbSSR.  
Predstavleno akademikom AN Azerbaydzhanskoy SSR I.D. Mustafayevym.  
(Bollworm)  
(Insecticides)

MAMDICOV, Shamkhal ~~MAZLOV~~, O.B.; KHYDYROV, D.N.; AVANECYAN, M.A.;  
AGAYEV, A.S.; GRISHINA, Ye.M.

The new contact insecticides efiran-79 and efiran-103 for  
agricultural pests. Dokl. AN Azerb. SSR 17 no.10:937-940  
'61. (MIRA 14:12)

1. Institut neftekhimicheskikh protsessov AN AzerSSR.  
Predstavleno akademikom AN AzSSR G.A. Aliyevym.  
(Insecticides)

MAMEDOV, Shamhal; ALIYEVA, Kh.M.; OSIPOV, O.B.

Utilization of olefinic components from the liquid pyrolysis  
products of petroleum hydrocarbons. Neftekhimia 2 no.1:115-  
120 Ja-F '62. (MDA 15:5)

1. Institut neftekhimicheskikh protsessov AN AzSSR.  
(Olefins) (Herbicides)

MAMEDOV, Shamkhal; OSIPOV, O.B.; DZHALILOV, T.N.; GRIZHINA, Ye.N.

New contact poisonous chemicals "efiran-168" and "efiran" 169."  
Dokl. AN Azerb. SSR 18 no.9:19-23 '62. (MIRA 17:1)

1. Institut neftekhimicheskikh protsessov AN AzerSSR.  
Predstavлено академиком AN AzSSR I.D. Mustafayevym.

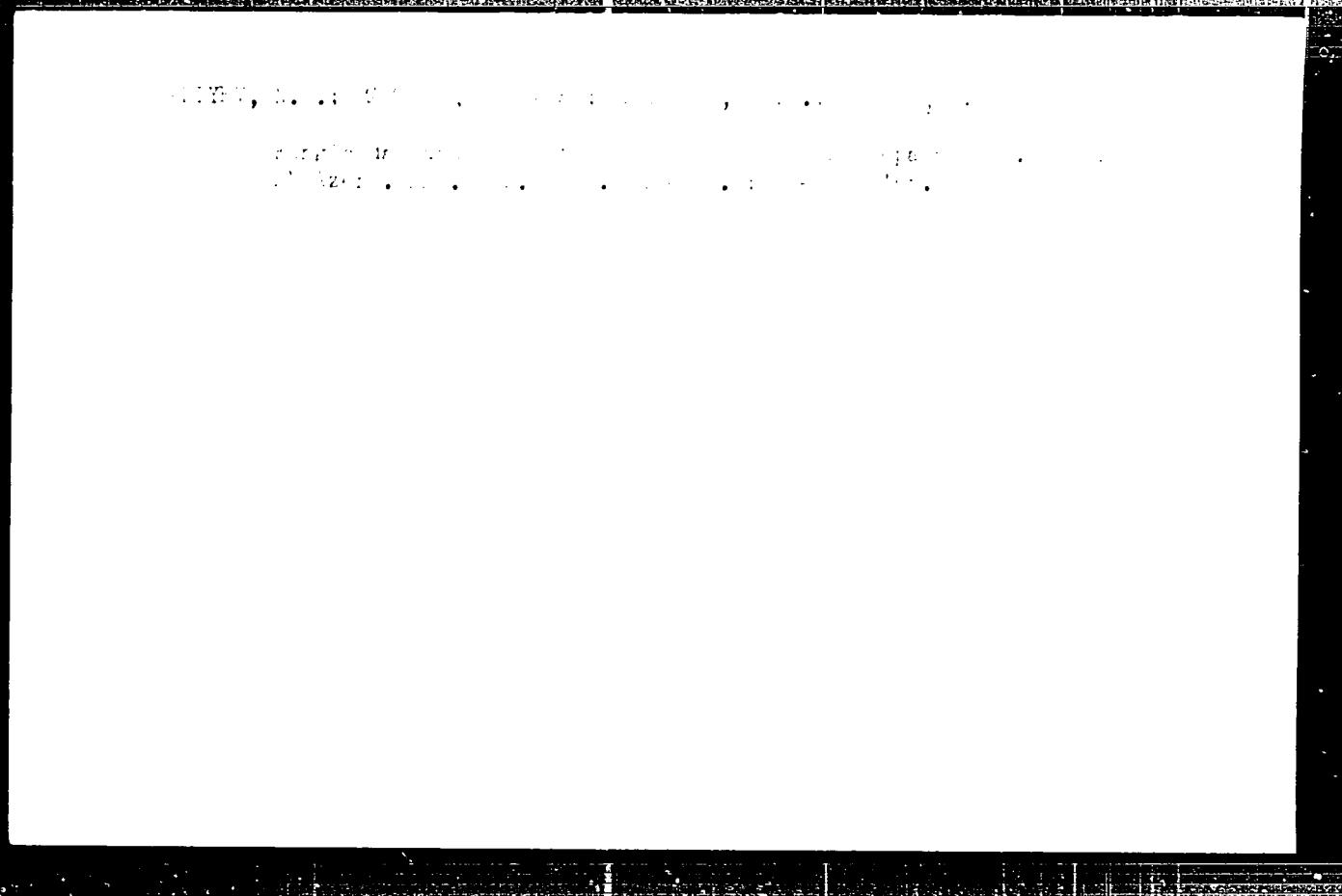
KESSLER, Yu.M.; ALPATOV, N.M., OJIPCOV, G.R.

Electrochemical and physicochemical properties of aluminum compounds  
in nonaqueous solutions. Vop. katal. 1976, no. 1-261-245. Mr. 164.

(MIRA 17-4)

1. Institut elektrokhimii AN SSSR.

"APPROVED FOR RELEASE: Wednesday, June 21, 2000 CIA-RDP86-00513R001238



APPROVED FOR RELEASE: Wednesday, June 21, 2000 CIA-RDP86-00513R001238

L 64382-63  
(4)

ACCESSION NR: AP5021616

UR/0286/65/000/013/0094/0094

11

B

AUTHORS: Mamedov, Sh.; Arabov, A. K. o.; Osipov, O. B.; Grishina, Ye. N.

TITLE: A method for controlling insects harmful to farm vegetation. Class 45,  
No. 172575

SOURCE: Byulleten' izobreteniy i tovarnykh znakov, no. 13, 1965, 94

TOPIC TAGS: pesticide, agriculture, insect control, organic compound

ABSTRACT: This Author Certificate presents a method for controlling insects harmful to farm vegetation by using an insecticide acting as a reaction starter. To enlarge the assortment of insecticides,  $\beta$ -diethylaminoethanol benzyl ether is used as a starting compound.

ASSOCIATION: none

SUBMITTED: 18Apr64

ENCL: 00

SUB CODE: LS, OG

NO REF Sov: 000

OTHER: 000

Cord 1/1 lla

KESSLER, Yu.M.; OSIPOV, G.R.

Electromagnetic chopper with a wide regulation range of the  
operational pulses. Zhur.fiz.khim. 39 no.11:2847-2848 N  
'65. (MIFKA 18:1.)

1. Institut elektrokhimii AN SSSR.

MOTORIN, I.I.; OSIPOV, O.V.; ARIFULIN, S.A.; PETRYAYEV, A.A., otv.  
red.; CHERNEGOVA, E.N., red. izd-va; LOMILINA, I.N.,  
tekhn. red.

[Regulations for the exploitation of drilling, pumping,  
compressor, power, special, and construction equipment]  
Pravila ekspluatatsii burovogo, nasosnogo, kompressornogo,  
silovogo, spetsial'nogo i stroitel'nogo oborudovaniia. Mo-  
skva, Gosgortekhizdat, 1962. 22 p. (MIRA 16:7)

1. Soyuzshakhtoosusheniye, trust.  
(Machinery)

L 52170-65	EWT(1)/EWA(3)/EWA(b)-2	Pa-4	RO
ACCESSION NR:	AP5015539	UR/0286/65/000/008/0080/0080	
AUTHORS:	Mamedov, Sh.; Mamedova, A. Kh. A.-K.; Avanesyan, M. A.; Osipov, O. V.; Zeynalova, V. M.; Karakhanova, S. V.		
TITLE:	A method for controlling weeds. Class 45, No. 170246 24 B		
SOURCE:	Byulleten' izobreteniya i tovarnykh znakov, no. 8, 1965, 80		
TOPIC TAGS:	agriculture, pesticide, ester		
ABSTRACT:	This Author Certificate presents a method for controlling weeds by herbicides. To broaden the assortment of <u>herbicides</u> , para- $\beta$ -cresoxy- $\beta$ -carbon-methoxydiethyl ester is used for this purpose.		
ASSOCIATION:	none		
SUBMITTED:	31Jul64	ENCL:	00
NO REP SOV:	000	OTHER:	000
zah Card 1/1			

"APPROVED FOR RELEASE: Wednesday, June 21, 2000

CIA-RDP86-00513R001238

OSIPOV, P.

All-purpose lathe for young technicians. IUn.tekh. 7 no.11:70-  
71 N '62.  
(Lathes) (MIRA 15:12)

APPROVED FOR RELEASE: Wednesday, June 21, 2000

CIA-RDP86-00513R001238

OSIPOV, P., nachal'nik; ZUBKOV, G., slesar'.

Window scaffold. Zhil.-kom. blok. 1 no.6:1R-21 Je 13. Matus

1. Remontno-stroitel'naya kontora Kirovskogo rayzhalupravleniya Rastov-na-Donu.  
(Scaffolding)

OSIPOV, P.

More about the lag in production of equipment for concrete work.  
Mekh. trud. rab. 9 no.7:18-19 Jl '55. (MLRA 8:9)

1. Nachal'nik Glavnogo upravleniya stroitel'nykh mashin i oborudovaniya dlya proizvodstva stroitel'nykh materialov Ministerstva stroitel'nogo i dorozhnogo mashinostroyeniya SSSR  
(Building machinery)

OSIPOV, P.A.

Causes of the formation of twist in raw silk during the unwinding  
of cocoons. Izv.vys.ucheb.zav.; tekhn.tekst.prom. no.2135-39 '63.  
(MIRA 16:6)

1. Kostromskoy tekhnologicheskiy institut.  
(Silk manufacture) (Winding machines)

BA

826-3

amount of hydrostatic pressure on the absorption and flow of water  
into the system. P. A. Chugov (Zhur. priro., 1980, No. 8, 18-21).  
Silk cocoons were submerged at depths of 0-90 cm. H<sub>2</sub>O at various

temp., and rates of absorption and amossis of H<sub>2</sub>O absorbed were  
measured. The effects of an immediate second immersion at 90-cm.  
depth were studied. The absorption and flow rate increase with  
depth of first immersion. effects of the second immersion are  
irregular, but at a given depth the flow rate at the second immersion  
is less than at the first, the max. difference being at 90'. Max  
absorption in the first immersion is at 90' and @ 90-cm. depth.  
E. B. Uzakov

OSIPOV, P.A.

Forces acting in the re-rolling during the movement of the "Kostromskoy Izv.vys.ucheb.zav.; tekhn.tekst.r m. no.32.3-50" [unclear]

1. Kostromskoy tekhnicheskoy institut.

"APPROVED FOR RELEASE: Wednesday, June 21, 2000

CIA-RDP86-00513R001238

OSIPOV, P.A.

Measuring of raw silk filaments. Izv. vys. ucheb. zav., tekhn.  
teks. prom. no.6:28-33 '65. (MIRD 14:1)

I. Kostroma Tekhnologicheskiy Institut. Submitted November 14,  
1964.

APPROVED FOR RELEASE: Wednesday, June 21, 2000

CIA-RDP86-00513R001238

BATASH V, Nikolay Nikanovich; KURNEV, Yevgeniy Mikhaylovich;  
OSIELEV, Petr Fedotovich; IGREVKII, V.I., red.;  
ISAYEVA, V.V., ved. red.; YAKOVIEVA, Z.I., tekhn.red.

[Preparation, treatment, and cleaning of clay solutions;  
practices of petroleum workers in Kuybyshev Province]  
Prigotovlenie, obrabotka i očistka glinistykh rastvorov;  
opyt neftianikov Kuybyshevskoi oblasti. Moscow, Gosstop-  
tekhizdat, 1963. 80 p. (MIRA 16:11)  
(oil well drilling fluids)

24 4100

26445  
S/021/60/000/004/001/010  
D232/D305

AUTHOR: Osypov, P.M.

TITLE: The inertial motor dyad

PERIODICAL: Akademiya nauk Ukrayins'koyi RSR. Dopovidi, no. 4,  
1960, 419 - 425

TEXT: The inertial properties of a moving particle are given by a single scalar - the mass of the particle. However, for the inertial properties of a moving rigid body to be fully determined, the following ten integrals are necessary: 1) The total mass

$$m = \int dm; \quad (1)$$

2) The three statical moments, giving the coordinates of the center of gravity

$$S_1 = mx_c = \int_V x dm, \quad S_2 = my_c = \int_V y dm, \quad S_3 = mz_c = \int_V z dm; \quad (2)$$

Card 1/5

26445  
S/021/60/000/004/001/010  
D232/D305

The inertial motor dyad

3) The three moments of inertia

$$I_{11} = \int_V (y^2 + z^2) dm, I_{22} = \int_V (z^2 + x^2) dm, I_{33} = \int_V (x^2 + y^2) dm; \quad (3)$$

4) The three products of inertia

$$I_{12} = I_{21} = - \int_V xy dm, I_{23} = I_{32} = - \int_V yz dm, I_{13} = I_{31} = - \int_V zx dm. \quad (4)$$

These 10 quantities are the components of a motor dyad which defines the combined inertial properties of the rigid body and which is called the inertial motor dyad. Theorem:

$$\hat{q} = mv + \hat{v} \cdot \hat{J} \quad (5)$$

where  $q$  is the motor of the quantity of motion,  $\hat{v} = \omega^* + \epsilon v$  is the velocity motor corresponding to the pole 0, and  $\hat{J}$  is the inertial motor dyad. It is shown that  $\hat{J} = J + J^*$ , where

Card 2/5

26445  
S/021/60/000/004/001/010  
D232/D305

The inertial motor dyad

$$J + \varepsilon J^* = \begin{bmatrix} 0 & S_3 - S_2 \\ -S_3 & 0 & S_1 \\ S_2 - S_1 & 0 \end{bmatrix} + \varepsilon \begin{bmatrix} I_{11} & I_{12} & I_{13} \\ I_{21} & I_{22} & I_{23} \\ I_{31} & I_{32} & I_{33} \end{bmatrix} \quad (13)$$

i.e.  $J$  is a hypercomplex dual number. Two orthogonal coordinate systems are chosen - a stationary system  $Oxyz$ , and a moving system  $Oxyz$ .  $r_o$  is the position-vector of the moving origin referred to the fixed axes, so that the position vector of a general point of the body is given by  $\bar{r} = r_o + r$ , where  $\bar{r}$  and  $r$  are the position vectors of the point referred to the stationary and moving systems respectively. The general equation of motion of a rigid body is discussed then. The derivative with respect to time of the motor of the quantity of motion equals the motor of the force applied to the body,  $k$ . Hence, from

$$\frac{d\hat{u}}{dt} = \frac{\partial \hat{u}}{\partial t} + \hat{v} \times \hat{u}, \quad (23)$$

Card 3/5

26445

The inertial motor dyad

S/021/60/000/004/001/010  
D232/D305

$$\frac{d\hat{q}}{dt} + \hat{v} \times \hat{q} = \hat{k}. \quad (27)$$

Applying the theorem, writing the expression in terms of  $J$  and simplifying gives

$$m \frac{\partial \hat{v}}{\partial t} + m\omega \times v + \frac{\partial \hat{v}}{\partial t} \cdot \hat{J} + \hat{v} \cdot \frac{d\hat{J}}{dt} = \hat{k}, \quad (34)$$

where

$$\frac{d\hat{J}}{dt} = \begin{bmatrix} 0 & \dot{s}_3 - \dot{s}_2 \\ -\dot{s}_3 & 0 & \dot{s}_1 \\ \dot{s}_2 - s_1 & 0 \end{bmatrix} + \varepsilon \begin{bmatrix} i_{11} & i_{12} & i_{13} \\ i_{21} & i_{22} & i_{23} \\ i_{31} & i_{32} & i_{33} \end{bmatrix} \quad (35)$$

Hence, the kinetic energy of the body is calculated as the dual part of the scalar product of the velocity motor by the motor of the quantity of motion of the rigid body

$$2T + mv^2 + \omega \cdot (\omega \cdot J^*) + \omega \cdot (v \cdot J) - v \cdot (\omega \cdot J). \quad (36)$$

Card 4/5

The inertial motor dyad

26445  
S/021/60/000/004/001/01  
D232/D305

There are 6 references: 4 Soviet-bloc and 2 non-Soviet-bloc.

ASSOCIATION: Kyyiv's'kyy tekhnolohichnyy instytut kharchovoy promy-slovosti (Kyyiv Technological Institute of the Food Industry)

PRESENTED: by Academician AS UkrSSR, O.Yu. Ishlins'kiy

SUBMITTED: May 5, 1959

✓

Card 5/5

*CIA RDP86*

24.420

28678  
S/021/60/000/007/001/009  
D211/D305

AUTHOR: Osypov, P.M.

TITLE: The motor diad of infinitesimal transformations

PERIODICAL: Akademiya nauk Ukrayins'koyi RSR. Dopovidi, no. 7,  
1960, 863 - 868

TEXT: The question of the continuous transformation of coordinates has an important theoretical as well as practical significance. The author aims at improving the formulae concerning this matter by R. Mises (Ref. 1: Zs. für ang. Math. u. Mech., 4, 3, 193, 1924) Mises' diad contains four times as many scalar components as the diad the author considers which has dual components. For  $t = 0$  the new coordinate system  $Ox'_1, x'_2 x'_3$  coincides with the old coordinate system  $Ox_1 x_2 x_3$ . For any time  $t$  its motion could be described by  $\dot{v}(v_1, v_2, v_3) = \omega + \epsilon \bar{v}(v_1, v_2, v_3)$ , i.e. the new origin  $O'$  moves in space with velocity  $v(v_1, v_2, v_3)$  and rotates round  $O'$  with angular velocity  $\omega$ .

Card 1/5

28678

S/021/60/000/007/001/009  
D211/D305

The motor diad of infinitesimal

lar velocity  $\omega(\omega_1, \omega_2, \omega_3)$ . For the diad of infinitesimal transformation the author uses a formula (21) from his previous paper (Ref. 2: DAS, Ukr RSR, 7, 1960)

$$U_\lambda = \sum_{x=1}^3 U_x i_x \cdot (i'_\lambda + \varepsilon r \cdot i_\lambda), \quad (\lambda = 1, 2, 3) \quad (2)$$

where

$$i'_\lambda = i_\lambda + \frac{di_\lambda}{dt} dt \quad (3)$$

and

$$\left. \begin{aligned} \frac{di_1}{dt} &= \omega \times i_1 = (\omega_1 i_1 + \omega_2 i_2 + \omega_3 i_3) \times i_1 = \omega_3 i_3 - \omega_2 i_1 \\ \frac{di_2}{dt} &= \omega_1 i_1 - \omega_3 i_3, \quad \frac{di_3}{dt} = \omega_2 i_2 - \omega_1 i_1 \end{aligned} \right\} \quad (4)$$

Applying formulae (3) and (2) and omitting the expressions with  $dt^2$  one obtains

Card 2/5

28678

8/021/60/000/007/001/009  
D211/D305

The motor diad of infinitesimal ...

$$\left. \begin{aligned} U_1' &= U_1(1 + \epsilon O) + U_2(-\omega_3 + \epsilon \omega_3)dt + U_3(-\omega_2 - \epsilon \omega_2)dt \\ U_2' &= U_1(-\omega_2 - \epsilon \omega_2)dt + U_2(1 + \epsilon O) + U_3(\omega_1 + \epsilon \omega_1)dt \\ U_3' &= U_1(\omega_1 + \epsilon \omega_1)dt + U_2(-\omega_1 - \epsilon \omega_1)dt + U_3(1 + \epsilon O) \end{aligned} \right\} \quad (7)$$

Therefore the matrix for the motor-diad of infinitesimal transformation is

$$\hat{\mathbf{A}} = \mathbf{A} + \epsilon \mathbf{A}^* = \begin{bmatrix} 1 + \epsilon O & V_3 dt - V_2 dt \\ -V_3 dt & 1 + \epsilon O & V_1 dt \\ V_2 dt & -V_1 dt & 1 + \epsilon O \end{bmatrix}. \quad (8)$$

Introducing the identical diad E and dividing by dt the author obtains

$$\hat{\mathbf{A}} = \hat{\mathbf{V}} dt + \mathbf{E} \quad \hat{\mathbf{A}}^c = \hat{\mathbf{V}}^c dt + \mathbf{E} = -\hat{\mathbf{V}} dt + \mathbf{E}. \quad (11)$$

Using Theorem (1) (Ref. 2: Op.cit.) and Eq. (11) a new diad is obtained ( $\hat{\mathbf{D}}'$ ) with respect to a new coordinate system.

Card 3/5

28678

S/021/60/000/007/001/009  
D211/D305

The motor diad of infinitesimal ...

$$\hat{D}' = \hat{A} \cdot \hat{D} \cdot \hat{A}^C = (\hat{V} dt + E) \cdot \hat{D} \cdot (-\hat{V} dt + E) \quad (12)$$

$$\frac{d\hat{D}}{dt} = \lim_{\Delta t \rightarrow 0} \frac{\hat{D}' - \hat{D}}{\Delta t} = \hat{V} \cdot \hat{D} - \hat{D} \cdot \hat{V}. \quad (13)$$

The author proves later that the derivative of the motor diad could be expressed as follows:

$$\frac{d\hat{D}}{dt} = \hat{D} \times \hat{V} - \hat{V} \times \hat{D} \quad (15)$$

and that the second member of the formula

$$\frac{d\hat{u}}{dt} = \frac{\hat{u}}{t} + \hat{v} \times \hat{u} \quad (21)$$

satisfies  $\hat{v} \cdot \hat{u} = \hat{u} \cdot \hat{v}$ . Applying his formula to the diad of inertia

ard4/5

28678

The motor diad of infinitesimal ...

S/021/60/000/007/001/009  
D211/D305

$$\mathbf{J} + \epsilon \mathbf{J}^* = \begin{bmatrix} 0 & S_2 - S_3 \\ -S_2 & 0 & S_1 \\ S_3 & -S_1 & 0 \end{bmatrix} + \epsilon \begin{bmatrix} I_{11} & I_{12} & I_{13} \\ I_{21} & I_{22} & I_{23} \\ I_{31} & I_{32} & I_{33} \end{bmatrix}. \quad (23)$$

he obtains

$$\frac{d\hat{\mathbf{J}}}{dt} = \dot{\mathbf{J}} + \epsilon \dot{\mathbf{J}}^* = \begin{bmatrix} 0 & S_2 - \dot{S}_3 \\ -\dot{S}_2 & 0 & \dot{S}_1 \\ S_3 - \dot{S}_1 & 0 & 0 \end{bmatrix} + \epsilon \begin{bmatrix} \dot{I}_{11} & \dot{I}_{12} & \dot{I}_{13} \\ \dot{I}_{21} & \dot{I}_{22} & \dot{I}_{23} \\ \dot{I}_{31} & \dot{I}_{32} & \dot{I}_{33} \end{bmatrix}. \quad (28)$$

which is important for the general theory of the motion of rigid bodies. There are 4 references: 3 Soviet-bloc and 1 non-Soviet-bloc.

ASSOCIATION: Kyyivs'kyy tekhnolohichnyy instytut kharchovoyi promyslovost (Kyyiv Technological Institute for the Food Industry)

PRESENTED: by O.Yu. Ishlins'kyy Academician AS UkrSSR

SUBMITTED: May 5, 1959

Card 5/5

S/021/60/000/008/004/011  
D210/D305

AUTHOR: Oseypov, P.M.

TITLE: Ostrogradsky's theorem in motor calculus

PERIODICAL: Akademiya nauk Ukrayins'koyi RSR. Dopovidi, no. 8,  
1960, 1019 - 1023

TEXT: The aim of the paper is to prove the formulae of R. Mises  
(Ref. 4: Zsch. fur ang. Math. u. mech., 4, 193, 1924) given without  
the proof. The motor is said to be any bivector received by  
reducing the systems of transmissible vectors to any point in the  
space. The motor-field is this part of the space which to each  
point some motor is prescribed respectively. The main argument  
the author uses is the proof of Ostrogradsky's theorem for the  
motor field and the motor diad. It has the following form:

$$\oint \mathbf{A} \cdot d\mathbf{s} = \int \nabla \cdot \mathbf{A} d\mathbf{s} \quad (4)$$

Card 1/5

Ostrogradsky's theorem in ...

S/021/60/000 '008/004/011  
D210/D305

or

$$\oint \mathbf{A}_n ds = \int \left( \frac{\partial A_1}{\partial x_1} + \frac{\partial A_2}{\partial x_2} + \frac{\partial A_3}{\partial x_3} \right) dt, \quad (5)$$

For the motor diad

$$\oint \mathbf{D} \cdot d\mathbf{s} = I_1 \int \nabla \cdot \hat{\mathbf{a}}_1 dt + I_2 \int \nabla \cdot \hat{\mathbf{a}}_2 dt + I_3 \int \nabla \cdot \hat{\mathbf{a}}_3 dt = \int \nabla \cdot (I_1 \hat{\mathbf{a}}_1 + I_2 \hat{\mathbf{a}}_2 + I_3 \hat{\mathbf{a}}_3) dt, \quad (7)$$

$$\oint \mathbf{D} \cdot d\mathbf{s} = \int \nabla \cdot \mathbf{D} dt.$$

where  $\hat{\mathbf{D}} = I_1 \hat{\mathbf{a}}_1 + I_2 \hat{\mathbf{a}}_2 + I_3 \hat{\mathbf{a}}_3$  is the motor diad field and  $\hat{\mathbf{a}}_1 = \mathbf{a}_1 - \epsilon \mathbf{a}^*$ . To find the motion of an elastic body, the author assumes agreement with R. Mises that in each point of a continuous medium act the resultant force motors  $\mathbf{K} = \mathbf{K} + \epsilon \mathbf{K}^*$  as well as force motors  $\hat{\mathbf{p}} = \mathbf{p} + \epsilon \mathbf{p}^*$  with respect to the unit surface. The generalized Newton's law for the continuous medium has the form

Card 2/5

Ostrogradsky's theorem in ...

S/021/60/000/008/004/011  
D210/D305

$$\frac{d\hat{q}}{dt} = \hat{k} + \hat{p} \quad (8)$$

where  $\hat{q}$  is the motor of linear momentum, and  $d\hat{q}/dt = \mu\hat{w}$  where  $\hat{w}$  is the motor of acceleration. Using formula Eq. (8) the equations of motion have the form

$$\int_{\tau_1}^{\tau_2} \mu\hat{w} d\tau = \int_{\tau_1}^{\tau_2} \hat{k} d\tau + \int_{\tau_1}^{\tau_2} \hat{p}_n ds \quad (13)$$

but

$$\oint_S \hat{p}_n ds = \int_{\tau_1}^{\tau_2} \nabla \cdot \hat{P} d\tau \quad (15)$$

and therefore  $\int_{\tau_1}^{\tau_2} \mu\hat{w} d\tau = \int_{\tau_1}^{\tau_2} \hat{k} d\tau + \int_{\tau_1}^{\tau_2} \nabla \cdot \hat{P} d\tau \quad (16)$

Since it is true for any volume  $\tau$   $\mu\hat{w} = \hat{k} + \nabla \cdot \hat{P}$  (17). By substituting  $\hat{p}_i = p_i + \epsilon(p_i^* + r \cdot p_i)$  ( $i = 1, 2, 3$ ) and using Eq. (17) one

Card 3/5

Ostrogradsky's theorem in ...

S/021/60/000/008/004/011  
D210/D305

obtains

$$\mu w + \epsilon (r \times w) = k + \nabla \cdot P + \epsilon (k_1' + r \times (k + \nabla \cdot P) + \nabla \cdot P^* + (p_{33} - p_{32}) I_1 + (p_{31} - p_{13}) I_2 + (p_{12} - p_{21}) I_3), \quad (20)$$

By comparison, two equations were obtained

$$\mu w = k + \nabla \cdot P, \quad (21) \quad (21)$$

$$\mu (r \times w) = k_1' + r \times (k + \nabla \cdot P) + \nabla \cdot P^* + (p_{33} - p_{32}) I_1 + (p_{31} - p_{13}) I_2 + (p_{12} - p_{21}) I_3, \quad (22) \quad (22)$$

By substituting (21) into Eq. (22) follows

$$0 = k_1' + \nabla \cdot P^* + (p_{23} - p_{32}) I_1 + (p_{31} - p_{13}) I_2 + (p_{12} - p_{21}) I_3. \quad (23)$$

If one assumes as usual that  $k_1' = 0$ ,  $P^* = 0$  then

$$p_{12} = p_{21}, \quad p_{13} = p_{31}, \quad p_{23} = p_{32}, \quad (24)$$

Card 4/5